

## The *Arabidopsis* pentatricopeptide repeat protein PDM1 is associated with the intergenic sequence of *S11-rpoA* for *rpoA* monocistronic RNA cleavage

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The *PDM1* gene encodes a pentatricopeptide repeat protein of the PLS subfamily. It is essential for *rpoA* polycistronic processing in *Arabidopsis*. In this study, we performed functional analysis of PDM1 in *rpoA* monocistronic cleavage and chloroplast development. The *pdm1* mutants display an albino lethal phenotype with severe chloroplast development defects. When the construct of *PDM1* fused with the *GFP* gene was introduced into *Arabidopsis* protoplasts, the GFP signal was exclusively observed in chloroplasts. This shows that PDM1 is localized to chloroplast. In the wild type, the *rpoA* transcript of about 990 nt is processed. In *pdm1*, this transcript is absent. However, Western blot showed that the RpoA protein in *pdm1* is accumulated at a level approximately 1/3 of the wild type. This suggests that certain other transcripts processed from *L23-L2-S19-L22-S3-L16-L14-S8-L36-S11-rpoA* polycistronic precursor may be used as templates for protein translation. To determine whether PDM1 can bind to the *rpoA* pre-mRNA, we generated a transgenic *Arabidopsis* line with *PDM1* fused to *FLAG* tag that was capable of complementing the *pdm1-1* mutant phenotype. RNA immunoprecipitation analysis showed that PDM1 is associated with *S11-rpoA* intergenic sequence. This indicates that PDM1 may bind to the *S11-rpoA* intergenic region to perform *rpoA* processing.

### *Arabidopsis*, PDM1, RNA cleavage

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Chloroplasts are derived from endosymbiotic cyanobacteria [1]. Throughout the course of evolution, the chloroplast genome has retained its prokaryotic characteristics and taken on certain new characteristics of the eukaryotic genome. As in the genome of cyanobacteria, many of the chloroplast genes are organized in multiple transcription units that are transcribed into polycistronic pre-mRNAs, which are then mostly cleaved into monocistronic mRNAs [2]. However, the chloroplast genome has evolved elaborate posttranscriptional RNA processing steps as well as eukaryotes. These processing steps include RNA splicing, RNA editing and exonucleolytic trimming for the purpose of stabilization and translation.

Recently, pentatricopeptide repeat proteins (PPRs) were reported to be involved in these RNA processing steps in chloroplasts. The PPR proteins are characterized by a degenerate motif of 35 amino acids that can be repeated up to 30 times within a single protein. Approximately half of the PPR proteins in *Arabidopsis thaliana* display tandem arrays of canonical repeats (or P-repeats). These proteins are known as P-type PPRs. Other family members, known as PLS subfamily, contain not only the standard 35 amino acid repeats, but also a shorter (S-repeats) and a longer version (L-repeats). The PLS subfamily proteins are further divided into PLS, E, E+ and DYW subgroups based on the presence of additional domains at the C terminus [3]. Numerous PPRs play roles in splicing [4–6]. Some of the PPR proteins may have an effect on both RNA stability and translation [7,8]. Fifteen

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PPRs have been found to have RNA-editing functions; they serve 22 of the 34 known editing sites in *Arabidopsis* [9].

PPR proteins are also reported to be involved in monocistronic transcripts cleavage from polycistronic ones. They include HCF152 [5] and CRR2 [10] in *Arabidopsis* and CRP1 [11] in maize. HCF152 is essential for endonucleolytic cleavage between *psbH* and *petB*. It binds certain areas of the *petB* transcripts. CRP1 is necessary for the processing of the monocistronic *petD* mRNA from the polycistronic precursor and for the translation of the chloroplast *petA* transcripts. It exhibits significant structural similarity to HCF152. CRR2 functions in the intergenic processing of chloroplast *rps7-ndhB* dicistronic transcript. It is essential for *ndhB* translation.

Plastid genes in higher plants are transcribed by plastid-encoded polymerase (PEP) and nucleus-encoded polymerase (NEP). Most plastid genes are transcribed by plastid-encoded polymerase (PEP). RpoA is a core subunit of PEP. In *Arabidopsis thaliana*, the monocistronic *rpoA* transcript of approximately 990 nucleotides is processed from the *L23-L2-S19-L22-S3-L16-L14-S8-L36-S11-rpoA* polycistron. The previous investigation shows that the PPR protein PDM1 is involved in the processing of the *rpoA* pre-mRNA in *Arabidopsis thaliana* [12]. In *pdm1*, the *rpoA* transcript of 990nt was absent. However, the molecular mechanism for PDM1 to regulate the *rpoA* mRNA processing is not clear. In this study, we performed functional analysis of PDM1 in *rpoA* monocistronic cleavage and chloroplast development in *Arabidopsis*.

## 1 Materials and methods

### 1.1 Plant material and growth conditions

*Arabidopsis* mutant *pdm1-4* (*Landsberg erecta* background) was screened using an EMS mutagenesis strategy. Prior to phenotypic analysis, *pdm1-4* was backcrossed to *Landsberg erecta* ecotype three times. The T-DNA insertion lines (*Columbia* ecotype) were obtained from the Arabidopsis Biological Resource Center (ABRC; Ohio State University). Seeds were sown on vermiculite and MS medium agar plates according to the different experimental procedures after imbibition for 3 d at 4°C. Plants were grown under long-day conditions (16 h of light/8 h of dark) at a constant temperature of 22°C.

### 1.2 Phenotype characterization and microscopy

Plants were photographed with a Nikon digital camera, Coolpix 4500 (Nikon, Japan). Small leaf segments were obtained from 2-week-old plants grown on MS medium agar plates added 2% sucrose. Transmission electron micrographs (TEMs) were obtained exactly as described [13]. The specimens were examined under transmission electron microscopy (Hitachi H7650; Hitachi, <http://www.hitachi.com>).

### 1.3 Molecular cloning of the PDM1 gene

The presence of the T-DNA insertion in the mutant was validated using primers that specifically amplified a sequence of the T-DNA (F: 5'-AGATTATTGCTCGGGTAG-AT-3'; R: 5'-GCTAATGGTAATGGTGCTAC-3'). Co-segregation of the T-DNA insertion site and the mutant phenotype were analyzed with AtLBb1.3 and the plant-specific primers (*pdm1-1*-LP: 5'-AAAGGCCATAAACTTGGAA-GC-3'; *pdm1-1*-RP: 5'-TTCATGGACCGCTATGATCTC-3'; *pdm1-2*-LP: 5'-TTCTTCTTTGCAATGGAATGG-3'; *pdm1-2*-RP: 5'-TGCACGTAAAGTGTTCGACAG-3'; *pdm1-3*-LP: 5'-TGGAAATGTGTCACTAAGGCC-3'; *pdm1-3*-RP: 5'-AAACACGGTCACATGGACTTC-3'). For each of the mutant plant, PCR using AtLBb1.3 and specific RP primers amplified a specific DNA fragment. For the WT plants, PCR using paired LP and RP primers amplified longer specific DNA fragments. PCR with both primer pairs exhibited positive results for the heterozygous mutant plants.

Complementation experiment was performed as follows: a DNA fragment of 3834 bp, including an 1983 bp upstream and 1851 bp coding sequence without a termination codon of *PDM1* was amplified and fused to a *FLAG*-tagged sequence (DYKDDDDK, nucleotides GACTACAAAGACGATGAC-GACAAG) using KOD plus polymerase (TAKARA, Japan) (CMP-F: 5'-CCATGGATACTGCTTCCAGTCATC-3'; CMP-R: 5'-AGATGTCTCCAAGTAACAAGTTGC-3'). Following sequence verification, the fragment was cloned into the pCambia1300 binary vector (Cambia; <http://www.cambia.org.au>), then the recombinant was introduced into heterozygous plants using the infiltration method by Agrobacterium strain LBA4404. The transformants were selected on PNS culture medium with 80 mg L<sup>-1</sup> hygromycin, and screened for plants with a green phenotype and a homozygous background. The primer sets were used as follows: AtLB3/RP primers were used to validate the presence of a T-DNA insertion in *pdm1-1*; genomic specific primers CI-F/CI-R (CI-F: 5'-TAGAAACACGGTCACATGG-3'; CI-R: 5'-ATTTGGTGCTATTGTGGAA-3') were used to validate the homozygous (*pdm1/pdm1*) background.

### 1.4 Subcellular localization of the PDM1 fused GFP protein

For GFP fusion, the 201-bp sequence containing the plastid transit peptide encoding sequence and the full-length CDS lacking a stop codon were respectively cloned from the seedling cDNA of the wild type with the following primers: *PDM1-TP*-F: 5'-ATGTTTTTCGTTATCGTTAAT-3'; *PDM1-TP*-R: 5'-TCCTTTGAAACCACTAAAAC-3'; *PDM1-CDS*-F: 5'-ATGTTTTTCGTTATCGTTAAT-3'; *PDM1-CDS*-R: 5'-AGATGTCTCCAAGTAACAAG-3'. The two sequences respectively fused with *eGFP* were cloned into the pMON530 binary vector. Transformation was performed as described above and the transformants were selected using 50 mg/L

kanamycin. The GFP fluorescence of the transgenic plants was observed under Confocal Laser Scanning Microscopy (LSM 5 PASCAL; ZEISS, <http://www.zeiss.com>).

### 1.5 Phylogenetic analysis

The multiple sequence alignment of the full-length protein sequences was performed using the ClustalX tool. The phylogenetic tree was constructed and tested by MEGA3.1 based on the UPGMA method.

### 1.6 Western blot analysis

Total leaf proteins were extracted, fractionated by SDS-PAGE and analyzed using immunoblotting, as described previously [14]. The anti-FLAG M2 monoclonal antibody was purchased from Sigma-Aldrich (<http://www.sigmaaldrich.com>), and the anti-RpoA polyclone antibody was raised against a synthetic peptide of RpoA by the GL Biochem company (<http://www.glbiochem.com>). The RpoA peptide was composed of the following amino acids: KCVESKRD-SKRLYYGR. We identified the activity and specificity of the anti-RpoA antibodies as follows: The *rpoA* gene from *Arabidopsis* was recombined with pET-51b vector and expressed in *E. coli* Rosseta strain. Western blot was then performed to verify the anti-RpoA antibodies against the expressing RpoA peptide in *E. coli* Rosseta strain. The empty pET-51b was used as a negative control.

### 1.7 RNA immunoprecipitation

*Arabidopsis* seedlings (21 d old) were fixed with 1% formaldehyde. Then the chloroplast isolation was performed according to Ketcham et al. [15]. The chloroplast was used for immunoprecipitation of specific RNA-protein complexes according to the description of Terzil et al. [16]. The anti-FLAG M2 monoclonal antibody and Dynabeads® ProteinG were purchased from Sigma-Aldrich and Invitrogen respectively (Sigma, <http://www.sigmaaldrich.com>; Invitrogen, <http://www.invitrogen.com>). RNA was isolated, purified and reverse-transcribed according to Cao et al. [17]. For real-time PCR, 1 µL of the cDNA was analyzed in a 20 µL reaction using the Quantitect SybrGreen PCR kit (Qiagen, <http://www.qiagen.com>) in an ABI7300 system (Bio-Rad, <http://www.bio-rad.com>). Each realtime PCR reaction was performed in triplicate. Primers used in the analysis are listed in Table S1.

## 2 Results

### 2.1 A point mutation in *PDMI* leads to an albino phenotype

To identify gene important for chloroplast development, an albino mutant was identified from a population of the *Ara-*

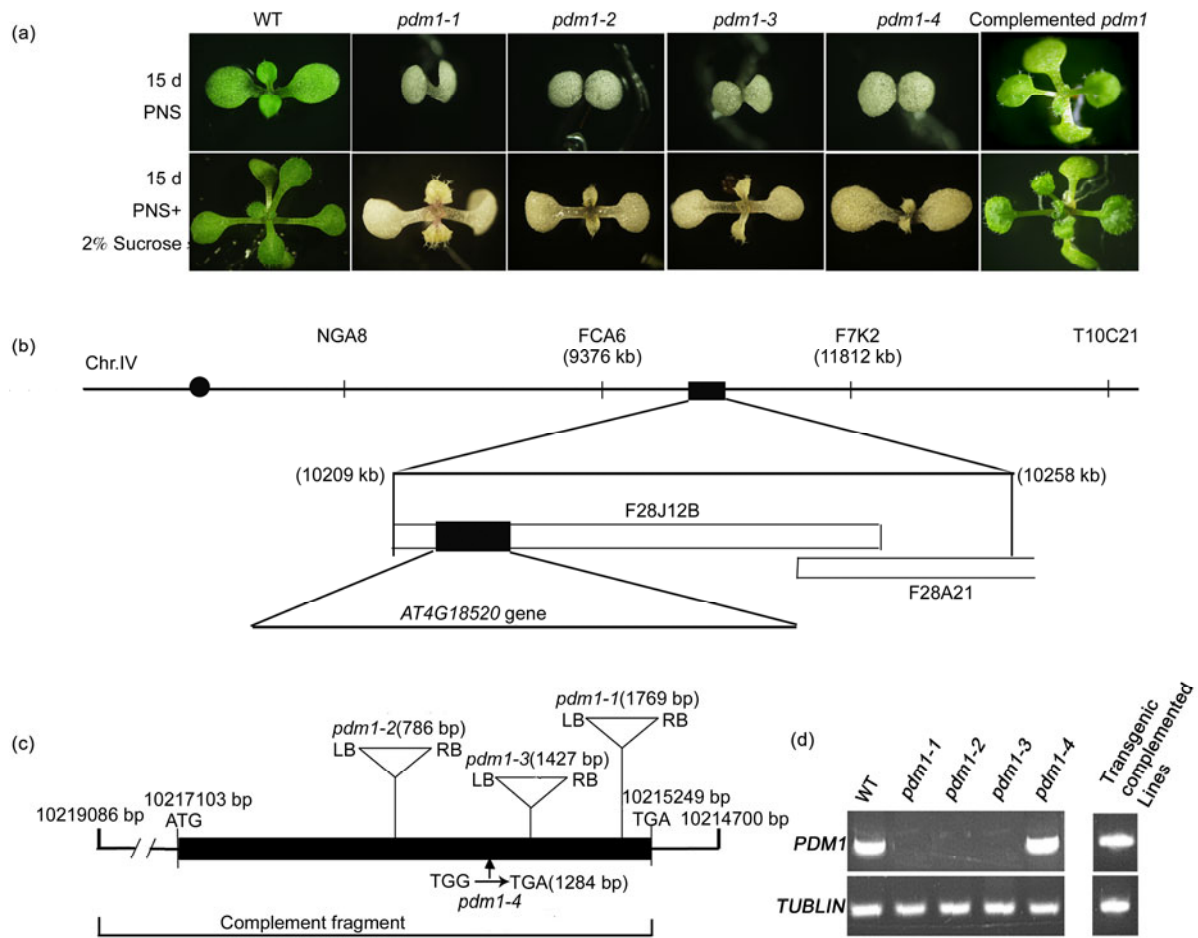
*bidopsis* Landsberg erecta ecotype (*Ler*) mutagenized with ethyl methane sulphonate (Figure 1(a)). Genetic analysis revealed this phenotype resulted from a single recessive locus (data not shown). The mutated locus was mapped using an F<sub>2</sub> population generated by crossing the heterozygote with the *Arabidopsis* Columbia ecotype (*Col*). For the first-pass mapping, a total of 20 insertion/deletion (In/Del) markers were used [18], and the mutated locus was linked to the In/Del marker FCA6 on chromosome 4. Fine mapping was carried out using a population of approximate 1800 albino progenies. The mutated locus was narrowed to a region of 49.6 kb between F28J12 and F28A21 on chromosome 4 (Figure 1(b)).

This region contains 13 putative genes, among which *PDMI* (*At4g18520*) encodes a PPR protein localized to chloroplast. Sequence analysis showed that a TGG to TGA mutation occurred in the mutant which caused translation to end prematurely in *PDMI*, resulting in a truncated form of 426 aa from a full length protein of 617 aa. We obtained three knockout lines for *PDMI* from the SIGnAL collection at the Arabidopsis Biological Resource Center (ABRC), *pdm1-1* (SALK\_034701), *pdm1-2* (CS835488) and *pdm1-3* (SALK\_005288). The T-DNA insertions in *PDMI* were confirmed by PCR analysis (Figure 1(c)). Semiquantitative RT-PCR analysis showed that the *PDMI* expression was not detectable in these mutants (Figure 1(d)). Allelism tests indicated that our mutant and these *pdm1* mutants belonged to the same locus mutation (data not shown). Therefore, our mutant is named *pdm1-4*. As the four alleles exhibit a similar phenotype, *pdm1-1* was chosen for further analysis.

To further investigate whether *PDMI* is responsible for the albino phenotype, we carried out a genetic complementation experiment. The *PDMI* genomic fragment, including a 1983-bp sequence upstream from the initiation codon, the 3' end of which lacks a termination codon, was fused to a FLAG-tagged sequence. This fused sequence was introduced into *PDMI/pdm1* heterozygous plants through *Agrobacterium tumefaciens*-mediated transformation. Eight transgenic plants carrying the complementation construct were obtained. These transgenic lines show the same phenotype as the WT. PCR analysis confirmed that the genetic background of one transgenic line was *pdm1/pdm1*. These results verified that *PDMI* was responsible for the *pdm1* albino phenotype.

### 2.2 Chloroplast development is defective in *pdm1-1* mutant

When grown in plant soil, the *pdm1-1* mutant displayed albino cotyledons. It could not survive more than 10 d after germination. When grown in MS medium containing 2% sucrose, it produced two to four etiolated true leaves. However, it died shortly after developing the second pair of true leaves (Figure 1(a)). To understand the chloroplast developmental defects in *pdm1-1*, the morphology and ultrastructure

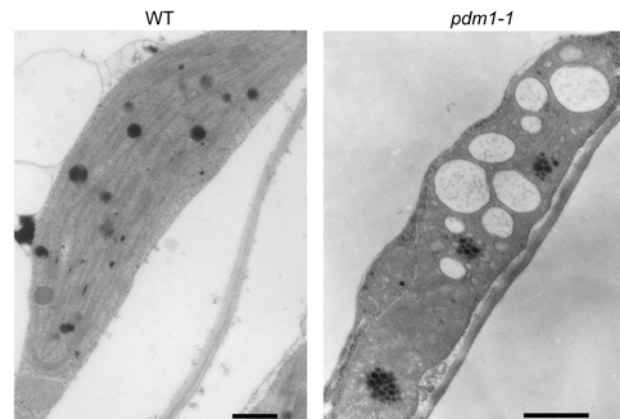


**Figure 1** Characterization of the *pdm1-1* mutant phenotype and the *PDM1* gene. (a) Phenotypes of the wild-type Columbia-0(WT), *pdm1-1*, *pdm1-2*, *pdm1-3* and *pdm1-4* mutants, and the complemented *pdm1*. (b) Mapping of the *PDM1* Locus. Black spot, the centromere. Fine mapping of *PDM1* to a 49.6-kb region between the In/Del markers F28J12 and F28A21 on chromosome 4. (c) The *PDM1* gene structure and T-DNA insertions of *pdm1-1*, *pdm1-2* and *pdm1-3*, and the position of the nucleotide change in *pdm1-4*. The complement fragment is indicated. (d) Expression analysis of the *PDM1* gene in the WT and *pdm1* mutants. *TUBULIN* expression is indicated as a control.

of the chloroplasts in 14-d-old leaves were examined under transmission electron microscopy (TEM). The mature WT chloroplasts exhibited normal stacked grana thylakoids, stroma thylakoids and starch grains. In contrast, only a few unstacked thylakoid lamella without stacked grana thylakoids were observed in the *pdm1-1* mutant (Figure 2). These results demonstrate that *PDM1* is essential for chloroplast development.

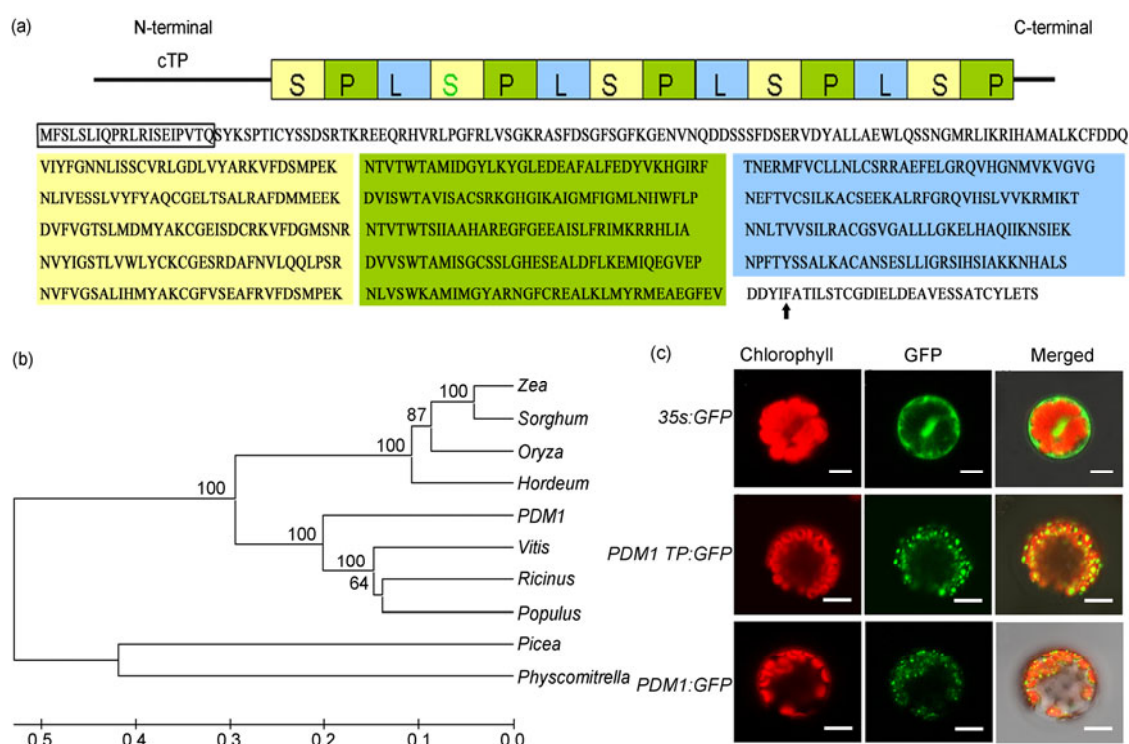
### 2.3 *PDM1* encodes a PPR protein localized to chloroplast

*Arabidopsis PDM1* encodes a PPR protein harbouring 14 PPR motifs, which are arranged as S-P-L-S-P-L-S-P-L-S-P-L-S-P. In the C terminus of the *PDM1* protein, there are extra 32 amino acids. However, this sequence is different from the currently identified E, E+ and DYW domains. Thus, *PDM1* is classified into the PLS subgroup. In the *pdm1-1* mutant, the last 28 amino acids are deleted, suggesting this region is essential for *PDM1* function (Figure 3(a)). Homologues of the *PDM1* protein have been identified



**Figure 2** Ultrastructure of chloroplasts in the wild-type and *pdm1-1*. In the mature chloroplasts of the WT, the stacked grana thylakoids and non-stacked stroma thylakoids are well organized, but in the *pdm1-1* mutant the early development of chloroplasts has been disturbed. Scale bars=0.5  $\mu$ m.

in various species by BLASTp or tBLASTn search in the NCBI database. Those proteins sharing a high degree of



**Figure 3** Phylogenetic analysis of PDM1 homologous proteins and subcellular localization of the PDM1 protein. (a) Protein structure of PDM1. The transit peptide is boxed. Black arrows indicate T-DNA interruption in the *pdm1-1* mutant. (b) Phylogenetic tree of PDM1 and homologous proteins. Protein sequence files: *Vitis vinifera*, XP\_002284293.1; *Ricinus communis*, XP\_002513855.1; *Populus trichocarpa*, XP\_002307403.1; *Hordeum vulgare*, BAJ99605.1; *Oryza sativa*, NP\_001173562.1; *Sorghum bicolor*, XP\_002464828.1; *Zea mays*, ACG29369.1; *Picea sitchensis*, ABR17838.1; *Physcomitrella*, PpPPR\_77. The amino acid sequences of the PDM1 homologous proteins were analyzed by the UPGMA method with the genetic distance calculated by Mega 3.1. (c) Subcellular localization of GFP fluorescence in the transformed protoplasts with the constructs of *PDM1::GFP* and *PDM1 TP::GFP*. Green fluorescence indicates the localization of PDM1::GFP and PDM1 TP::GFP proteins, while the red fluorescence is the autofluorescence of chlorophyll. Scale bars=20 μm.

similarity with the PDM1 are used for phylogenetic analysis. In the phylogenetic tree PDM1 forms a basal clade within the dicotyledons (Figure 3(b)). The results indicated that PDM1 has been conserved in the evolutionary process, and that putative homologues are present in various land plants, including mosses, gymnosperm and angiosperm.

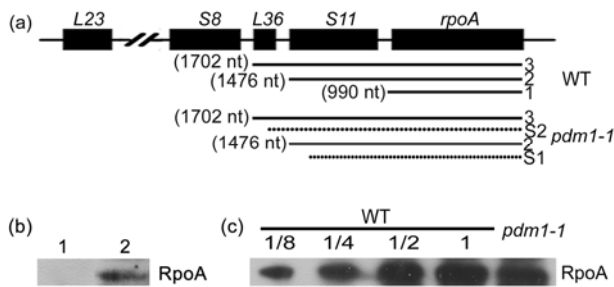
TargetP program analysis showed that the 19 amino acids at the N-terminus of PDM1 had the capacity to function as a plastid transit peptide (<http://www.cbs.dtu.dk/services/TargetP>) [19]. To investigate this possibility, we fused the first 67 amino acids and the full-length PDM1 protein to the N-terminus of a green fluorescent protein (GFP) to obtain *PDM1 TP::GFP* and *PDM1::GFP* constructs. These constructs, driven by a cauliflower mosaic virus (CaMV) 35S promoter, were transformed into *Arabidopsis* protoplasts. At the same time, the only GFP driven by the 35S promoter was transformed into protoplasts as a control. In the protoplasts transformed with the control construct, the GFP fluorescence was observed in the cytoplasm. However, in the protoplasts transformed with the *PDM1 TP::GFP* construct or *PDM1::GFP* construct, the GFP fluorescence co-localized with the red autofluorescence of chlorophyll (Figure 3(c)). These results confirm that the *Arabidopsis* PDM1 protein is localized to the chloroplast.

## 2.4 The defective processing of the *RpoA* polycistronic transcript leads to a low accumulation level of the *RpoA* protein

It was reported that monocistronic *rpoA* transcript cannot be detectable and that PEP-dependent chloroplast gene expression is affected in the *pdm1-1* mutant [12]. In the *pdm1-1* mutant, the 990 nt transcript for *rpoA* was lacking, while two new transcripts S1 and S2 were processed. The transcript S1 is shorter than the dicistronic transcript *S11-rpoA*. The transcript S2 is longer than the *S11-rpoA* dicistronic transcript, but shorter than the *L36-S11-rpoA* tricistronic transcript (Figure 4(a)).

To investigate whether the translation of RpoA was affected, a polyclonal antibody against RpoA was prepared in rabbits. A peptide specific for RpoA was used for immunization. The specific anti-RpoA antibody was purified, and then the RpoA antibody was tested for activity and specificity using recombinant protein (Figure 4(b)). This verified antibody was used to analyze the accumulation of RpoA protein in the *pdm1-1* mutant and the WT. The result showed that the amount of RpoA in the *pdm1-1* mutant is approximately 1/3 the level in the WT (Figure 4(c)). These results indicate that the defective *rpoA* polycistronic processing affects the



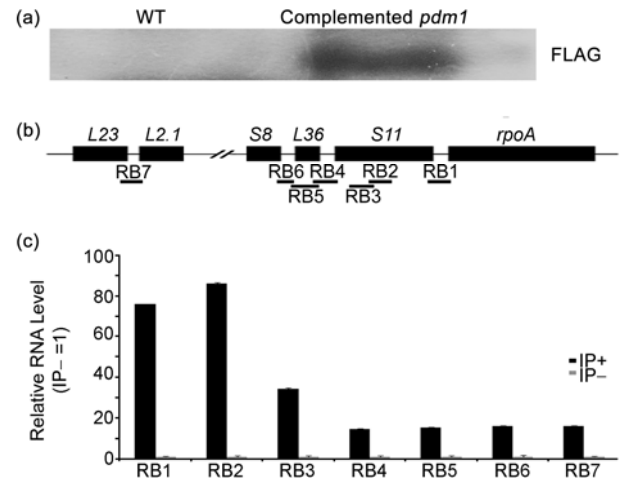


**Figure 4** Schematic diagram of the *rpoA* transcripts and the change in RpoA protein accumulation in the *pdm1-1* mutant. (a) Schematic diagram of smaller transcripts containing *rpoA* according to Wu et al. [12]. Black lines, normal transcripts in the WT and the *pdm1-1* mutant; dashed lines, additional transcripts in the *pdm1-1* mutant. (b) Activity and specificity identification of anti-RpoA antibody. The *rpoA* gene from *Arabidopsis* recombined with pET-51b vector was introduced into *E. coli* Rosseta strain. 1, Lysate of Rosseta expressing pET-51b; 2, lysate of Rosseta expressing pET-51b-RpoA. (c) RpoA protein accumulation characterization. Total proteins were extracted from 2-week-old WT and *pdm1-1* mutant plants, respectively, then size-fractionated by SDS-PAGE, transferred to a PVDF membrane, and detected with anti-RpoA antibodies.

translation and accumulation of the RpoA protein in the *pdm1-1* mutant.

## 2.5 PDM1 is associated with the *rpoA* polycistronic transcripts

PPR motifs have been demonstrated to have the capacity to bind RNA transcripts *in vivo* [6,8,20,21]. To determine whether PDM1 associate with *rpoA* pre-mRNA, RNA immunoprecipitation (RIP) was performed. In the complementation experiment, we obtained a transgenic *Arabidopsis* line that expresses PDM1 fused to the FLAG tag which can rescue the albino phenotype (Figure 1(a)). Expression of the fusion protein was verified by Western blot assay with an anti-FLAG M2 monoclonal antibody (Figure 5(a)). This transgenic line was used for RIP analysis. The PDM1-FLAG protein was immunoprecipitated from the crude chloroplast protein extract of the transgenic plants using Dynabeads decorated with anti-FLAG antibodies (IP+). As a negative control (IP-), Dynabeads without antibodies were used. Then RNA extracted from the immunoprecipitated products was used for quantitative real-time RT-PCR analysis. When the primer pairs for the intergenic region of *S11-rpoA* (designated RB1) and the *L23-L2.1* (designated RB7) were used (Figure 5(b)), the copy number of the PCR products from the anti-FLAG precipitate was nearly 88-fold and 14-fold that of those from the mock immunoprecipitation respectively (Figure 5(c)). To further understand the binding of PDM1 with *rpoA* pre-mRNA, we quantified the RNA level of the fragments RB2, RB3, RB4, RB5 and RB6 in the anti-FLAG precipitate (Figure 5(b)). The quantitative real-time RT-PCR results showed that RB2 in the anti-FLAG immunoprecipitated samples had a similar enrichment level as RB1, being nearly 88-fold that of mock



**Figure 5** RNA immunoprecipitation analysis of PDM1 and associated RNA. (a) Verification of PDM1-FLAG protein accumulation. Total proteins were extracted respectively from 2-week-old WT and complemented *PDM1-FLAG* transgenic plants, then immunoblot analysis was performed with anti-FLAG antibodies. (b) Distribution of the gene fragments in the polycistron *L23-L2-S19-L22-S3-L16-L14-S8-L36-S11-rpoA* for RNA immunoprecipitation analysis. (c) RNA immunoprecipitation analysis. Chloroplasts fixed with FA were extracted from the *PDM1-FLAG* transgenic plants, and then all of the RNA transcripts were sonicated into short fragments under specific conditions and incubated with a Dynabeads-linked FLAG antibody. With the interaction between FLAG antibody and PDM1-FLAG protein, associated RNA was immunoprecipitated and analyzed through realtime RT-PCR. IP+, anti-FLAG immunoprecipitation; IP-, mock immunoprecipitation. Data presented are mean values obtained from three replicates. Bars denote standard errors.

immunoprecipitation. However, RB3, RB4, RB5 and RB6 exhibited much lower enrichment (Figure 5(c)). These results suggest that PDM1 can bind to the *S11-rpoA* intergenic sequence.

## 3 Discussion

In this study, the function of the PPR protein PDM1 in RNA processing and chloroplast development was investigated in *Arabidopsis*. A defect in *rpoA* processing resulted in a low level of accumulation of RpoA protein, which exerted an effect on the expression of PEP-dependent plastid genes and chloroplast development in the knockout mutant *pdm1-1*. The RNA immunoprecipitation results suggest that PDM1 binds to the intergenic sequence between *rpoA* and *S11* to carry out endonucleolytic cleavage of *rpoA* from the precursor polycistronic transcripts.

### 3.1 PDM1 is essential for chloroplast development in *Arabidopsis thaliana*

Many factors have been reported to influence chloroplast development, including ribosome assembly [22], thylakoid membrane formation [23] and chloroplast gene expression [24]. In chloroplast development, PEP is indispensable for

chloroplast gene expression as part of the essential transcription machinery. In recent years, many proteins have been reported to regulate PEP activity, with most of them belonging to the plastid Transcriptionally Active Chromosome (pTAC) complex. The knockout of these genes affects PEP activity and chloroplast development [25,26]. According to the results presented here, the *pdm1-1* mutant displays an albino phenotype with severely defective chloroplast development (Figure 2), while the results of Wu et al. [12] reveals that PEP-dependent gene expression is dramatically reduced in this mutant. These suggest that the defective PEP activity in *pdm1* is the major factor that results in its lethal albino phenotype.

RpoA is a core subunit of PEP. The normal 990 nt *rpoA* transcript is cleaved from polycistronic transcripts and is responsible for efficient RpoA protein synthesis in the WT. In this work, our results show that PDM1 is associated with the *rpoA* polycistronic transcript (Figure 5). However, a new transcript of approximate 1300 nt (S1, Figure 4(a)) was present in the *pdm1-1* mutant instead of the 990 nt transcript observed in the WT [12]. The RpoA protein can be translated with low efficiency and its accumulation amounted to about 1/3 of the WT (Figure 4). This suggests other transcripts of *rpoA* may be used as templates for protein translation. This decreased RpoA amount in *pdm1* would be expected to exert an effect on PEP activity. However, PEP seems to have very low activity in *pdm1-1* mutant [12]. The PPR proteins have a range of essential functions in post-transcriptional processes [3]. For instance, PPR10 can stimulate *atpH* translation in addition to stabilizing the 5' and 3' chloroplast mRNA termini [8]. Additionally, the PPR protein CRP1 is required for the accumulation of processed transcripts of *petB* and *petD* in maize chloroplasts and the translation of RNAs [11,20]. Given the complexity of the functions of the PPR proteins, the possibility cannot be ruled out that PDM1 may be involved in an as yet unidentified process which affects PEP activity.

### 3.2 PDM1 is involved in the endonucleolytic cleavage of *S11* and *rpoA*

*PDM1* encodes a PPR protein of the PLS subfamily (Figure 3). PPR motifs have been demonstrated to have a capacity to bind RNA and the PDM1 protein consists of 14 PPR motifs. Since the RIP result showed that PDM1 was associated with the *S11-rpoA* intergenic sequence, PDM1 may bind to *rpoA* pre-mRNA. In the C terminus of PDM1, there is an extra 32 amino acid sequence that does not belong to PPR motif. In the *pdm1-1* mutant, 28 of the 32 amino acids are deleted, and two new transcripts are processed instead of the 990 nt transcript for *rpoA* (Figure 4(a)). This cleavage pattern change suggests that the C terminal region in PDM1 is responsible for the recognition of the PDM1 or PDM1 complex on the polycistronic precursor. However, there are no reports to show that PPR proteins have catalytic activity.

PDM1 might associate or interact with other proteins in order to perform the cleavage function. The amino acid sequence from Val<sup>116</sup> to Val<sup>585</sup> in PDM1 composes tandem arrays of PPR motifs. In the N terminus of the PDM1 protein, the first 19 amino acid sequence is predicted to be a transit peptide (Figure 3(a)). The amino acid sequence from Gln<sup>20</sup> to Gln<sup>115</sup> may play a role in the interaction with other proteins. The identification of proteins interacting with PDM1 should facilitate our understanding of the molecular mechanism that PDM1 regulate *rpoA* monocistronic RNA cleavage.

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## Supporting Information

**Table S1** Primers for RNA immunoprecipitation analysis

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